## **AMENDMENTS TO THE CLAIMS**

The below listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently amended) A method for detecting a neoplasia in a biologic sample, the method comprising quantifying the promoter methylation of at least two promoters in the sample, wherein one of the promoters is pi-class glutathione S-transferase (GSTP1) and the second promoter is selected from the group consisting of O<sup>6</sup> methylguanine DNA methyltransferase (MGMT), p14/ARF, p16/INK4a, RAS associated domain family 1A (RASSFIA), adenomatous polyposis coli (APC), a Ras-associated domain family 1A (RASSF1A) promoter, and a cellular retinoid binding protein 1 (CRBP1) promoter,

tissue inhibitor of metalloproteinase 3 (TIMP3), S100A2, cellular retinoid binding protein 1 (CRBP1), and retinoic acid receptor  $\beta$ 2 (RAR $\beta$ 2), wherein an increased quantity of promoter methylation relative to a reference indicates the presence of a neoplasia in the sample.

- 2. (Currently amended) The method of claim 1, wherein the <u>method further</u> comprises quantifying the promoter methylation of second promoter is selected from the group consisting of *APC*, *RASSF1A*, *CRBP1*, and  $RAR\beta2$ .
  - 3-4. (Canceled)
- 5. (Withdrawn) A method of determining the clinical aggressiveness of a neoplasia in a biologic sample, the method comprising quantifying the level of *GSTP1* or *APC* promoter methylation in the sample, wherein an increased level of promoter methylation relative to a reference indicates an increased clinical aggressiveness of the neoplasia.
- 6. (Withdrawn) A method of determining the stage of a neoplasia in a biologic sample, the method comprising quantifying the level of promoter methylation in the sample of at least one promoter selected from the group consisting of GSTP1, APC, RASSF1A, and  $RAR\beta2$ , wherein an increased level of promoter methylation in the sample relative to a reference indicates an increased stage of neoplasia.

Amendment dated September 14, 2011 Reply to Office Action of March 14, 2011

- 7. (Previously presented) The method of claim 1, wherein the neoplasia is prostate cancer.
  - 8-9. (Canceled).
- 10. (Withdrawn) The method of claim 6, wherein the sample is a prostate tissue sample.
- 11. (Withdrawn) The method of claim 6, wherein the sample is a biologic fluid selected from the group consisting of serum, plasma, ejaculate, or urine.
  - 12. (Canceled)
- 13. (Previously presented) The method of claim 1, wherein the reference is the level of methylation present at the promoter in a control sample derived from a healthy subject.
  - 14. (Canceled)
- 15. (Previously presented) The method of claim 1, wherein the promoter methylation is quantified by quantitative methylation-specific PCR (QMSP).
- 16. (Previously presented) The method of claim 15, wherein the level or the frequency of promoter methylation is quantified.
  - 17. (Canceled)
  - 18. (Canceled)
  - 19-36. (Canceled)
- 37. (Currently amended) A method of determining the prognosis of a subject diagnosed as having a neoplasia, the method comprising quantifying the level of promoter methylation in a sample derived from the subject, wherein at least one the promoters are is selected from the group consisting of *GSTP1*, *APC*, *RASSF1A*, and *CRBP1*, and *RARβ2*, and

wherein an altered level of promoter methylation relative to a reference indicates the prognosis of the subject.

- 38. (Previously presented) The method of claim 37, wherein the alteration is a decrease or an increase in the level of promoter methylation relative to a reference.
- 39. (Previously presented) The method of claim 38, wherein the increased or decreased level of promoter methylation indicates a poor prognosis or a good prognosis.

## 40-43. (Canceled)

- 44. (Currently amended) A method of monitoring a subject diagnosed as having a neoplasia, the method comprising quantifying the level of promoter methylation in a sample derived from the subject, wherein at least one promoter is selected from the group consisting of the promoters are GSTP1, APC, RASSF1A, and CRBP1, and  $RAR\beta2$ , and wherein an altered level of promoter methylation relative to the level of methylation in a reference indicates an altered severity of neoplasia in the subject.
- 45. (Previously presented) The method of claim 37, wherein the reference is the level of methylation present in a sample previously obtained from the subject prior to therapy or present in a normal patient sample.

## 46-50. (Canceled)

- 51. (Original) The method of claim 44, wherein the sample is a prostate tissue sample.
- 52. (Original) The method of claim 44, wherein the patient sample is a biologic fluid selected from the group consisting of serum, plasma, ejaculate, or urine.
  - 53. (Canceled)
- 54. (Previously presented) The method of claim 37, wherein the promoter methylation is quantified by quantitative methylation-specific PCR.

Amendment dated September 14, 2011 Reply to Office Action of March 14, 2011

55-57. (Canceled)

- 58. (Currently amended) A method of selecting a treatment for a subject diagnosed as having a neoplasia, the method comprising:
- (a) quantifying the level of promoter methylation in a biologic sample from the subject relative to a reference, wherein the promoters are *GSTP1*, *APC*, *RASSF1A*, and *CRBP1*, and the level of promoter methylation is indicative of a treatment; and
  - (b) selecting a treatment.
- 59. (Previously presented) The method of claim 58, wherein the neoplasia is prostate cancer.
- 60. (Withdrawn) A method of selecting a treatment for a subject diagnosed as having prostate cancer, the method comprising:
- (a) quantifying the level of promoter methylation of a promoter selected from the group consisting of GSTP1, APC, RASSF1A, CRBP1, and  $RAR\beta2$  in a subject sample; and
- (b) selecting a treatment for the subject, wherein the treatment is selected from the group consisting of surveillance, surgery, hormone therapy, chemotherapy, and radiotherapy.
- 61. (Original) A method for determining the methylation profile of a prostate cancer, the method comprising quantifying the level of promoter methylation at two or more promoters selected from the group consisting of GSTP1, APC, RASSF1A, CRBP1, and  $RAR\beta2$  in a biologic sample, wherein the level of promoter methylation relative to a reference determines the methylation profile of the prostatic neoplasia.
- 62. (Withdrawn) A kit for the analysis of promoter methylation, the kit comprising at least one primer capable of distinguishing between methylated and unmethylated promoter sequences, wherein the promoter sequences are selected from the group consisting of GSTP1, APC, RASSF1A, CRBP1, and  $RAR\beta2$ , and directions for using the primer for the analysis of promoter methylation.

63. (Withdrawn) the kit of claim 62, wherein at least one of the primers binds selectively to a methylated or unmethylated sequence.

## 64-68. (Canceled)

- 69. (Withdrawn) A microarray comprising at least two nucleic acid molecules, or fragments thereof, bound to a solid support, wherein the two nucleic acid molecules are selected from the group consisting of *GSTP1*, *MGMT*, *p14/ARF*, *p16/INK4a*, *APC*, *RASSF1A*, *TIMP3*, *S100A*, *CRBP1*, and *RARβ2*.
- 70. (Currently amended) A method for detecting a neoplasia in a biologic sample, the method comprising quantifying the promoter methylation of at least two promoters in the sample by contacting the sample with a microarray of claim 59, wherein one of the promoters is selected from the group consisting of are GSTP1, MGMT, p14/ARF, p16/INK4a, APC, RASSF1A, TIMP3, S100A, and CRBP1, and RAR $\beta$ 2, and wherein an increased quantity of promoter methylation relative to a reference indicates the presence of a neoplasia in the sample.
- 71. (Withdrawn) A primer having a nucleic acid sequence selected from the group consisting of:
  - 5'- TGG TTT CGA TTT TTT GAT TTC G -3' (SEQ ID NO: 12).
  - 5'- TCA AAA TTC TTT TTA CAA CAA CGC C -3' (SEQ ID NO: 13),
  - 5'- CTG GGA ATC CAG CTG TCG CCG CCC CGC A -3' (SEQ ID NO: 15),
  - 5'- GCG CAT CAT AGC CAT CAG CAA CAA A -3' (SEQ ID NO: 16),
  - 5'-CGA GAA CGC GAG CGA TTC-3' (SEQ ID NO: 18),
  - 5'-CAA ACT TAC TCG ACC AAT CCA ACC-3' (SEQ ID NO: 19),
  - 5'-TGG TGA TGG AGG AGG TTT AGT AAG T-3' (SEQ ID NO: 21)
  - 5'- AAC CAA TAA AAC CTA CTC CTC CCT TAA-3'(SEQ ID NO:22).

- 72. (Withdrawn) A probe having a nucleic acid sequence selected from the group consisting of:
  - 5'- CGA CCG AAC GCG ATA ACT TAC TCC -3'-TAMRA (SEQ ID NO:14),
  - 5'- GAC CCG AAA ATA AAC GCC CTC CGA AAA CA -3' (SEQ ID NO:17),
- 5'-TCG GAA CGT ATT CGG AAG GTT TTT TGT AAG TAT TT-3' (SEQ ID NO: 20),
  - 5'-ACC ACC ACC CAA CAC ACA ATA ACA AAC ACA-3' (SEQ ID NO: 23).
- 73. (Withdrawn) A collection of primer sets, each of said primer sets comprising at least two primers that bind to a promoter selected from the group consisting of *GSTP1*, *MGMT*, *p14/ARF*, *p16/INK4a*, *APC*, *RASSF1A*, *TIMP3*, *S100A*, *CRBP1*, and *RARβ2*, said collection comprising at least two primer sets.